## RECEIVED CENTRAL FAX CENTER

AUG 3 - 2006

PATENT

Appl. No. 10/789,159 Amdt. dated August 3, 2006 Reply to Office Action of July 3, 2006

## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

## Listing of Claims:

- 1. (Original) A method for inducing differentiation of pluripotent cells comprising the following steps (a) and (b):
  - (a) culturing the pluripotent cells in a medium comprising any one of the following growth factors (i) to (iii):
    - (i) acidic fibroblast growth factor, fibroblast growth factor 4, and hepatocyte growth factor;
    - (ii) acidic fibroblast growth factor, and growth factor(s) selected from activin A, epidermal growth factor, and  $\beta$ -nerve growth factor; and
    - (iii) fibroblast growth factor 4, and growth factor(s) selected from activin A and hepatocyte growth factor; and,
  - (b) culturing the cell cultured in step (a) in a medium comprising oncostatin M.
- 2. (Original) The method according to claim 1, wherein a gelatin-coated culture dish is used in step (a), and a collagen type I-coated culture dish or laminin-coated culture dish is used in step (b).
- 3. (Original) The method according to claim 1, wherein a collagen type I-coated culture dish is used.
- 4. (Original) A method for inducing differentiation of pluripotent cells comprising the following steps (a) and (b):
  - (a) culturing the pluripotent cells in a medium comprising at least one growth factor selected from retinoic acid, leukemia inhibitory factor, and hepatocyte growth factor; and,
  - (b) culturing the cell cultured in step (a) in a medium comprising any one of the following growth factors (i) to (iii):

Appl. No. 10/789,159 Amdt. dated August 3, 2006 Reply to Office Action of July 3, 2006 PATENT

- (i) acidic fibroblast growth factor, fibroblast growth factor 4, and hepatocyte growth factor;
- (ii) acidic fibroblast growth factor, and growth factor(s) selected from activin A, epidermal growth factor and  $\beta$ -nerve growth factor; and
- (iii) fibroblast growth factor 4, and growth factor(s) selected from activin A and hepatocyte growth factor.
- 5. (Original) The method according to claim 3, wherein gelatin-coated culture dishes are used in steps (a) and (b).
- 6. (Original) A method for inducing differentiation of pluripotent cells comprising the following steps (a) to (c):
  - (a) culturing the pluripotent cells in a medium comprising at least one of the growth factors selected from retinoic acid, leukemia inhibitory factor and hepatocyte growth factor;
  - (b) culturing the cell cultured in step (a) in a medium comprising any one of the following growth factors (i) to (iii):
    - (i) acidic fibroblast growth factor, fibroblast growth factor 4 and hepatocyte growth factor;
    - (ii) acidic fibroblast growth factor, and growth factor(s) selected from activin A, epidermal growth factor and  $\beta$ -nerve growth factor; and
    - (iii) fibroblast growth factor 4, and growth factor(s) selected from activin A and hepatocyte growth factor, and,
  - (c) culturing the cells cultured in step (b) in a medium comprising oncostatin M.
- 7. (Previously presented) The method according to claim 6, wherein gelatin coated culture dishes are used in steps (a) and (b), and a collagen type I-coated culture dish or laminin-coated culture dish is used in step (c).
- 8. (Previously presented) A method according to claim 1, wherein the pluripotent cells are derived from a mammal.

<u>PATENT</u>

Appl. No. 10/789,159 Amdt. dated August 3, 2006 Reply to Office Action of July 3, 2006

- 9. (Original) The method according to claim 8, wherein the mammal is a human, monkey, mouse, rat or pig.
- 10. (Previously presented) A method according to claim 1, wherein the pluripotent cells are embryonic stem cells, adult stem cells, mesenchymal stem cells, or umbilical cord blood cells.
- 11. (Previously presented) A method for producing hepatocytes, wherein the method comprises steps (a) and (b) according to claim 1, or steps (a) to (c) according to claim 6.
- 12. (Original) The method according to claim 11, wherein the hepatocytes are mature hepatocytes.
- 13. (Previously presented) The method according to claim 11, wherein the pluripotent cells are derived from a mammal.
- 14. (Original) The method according to claim 13, wherein the mammal is a human, monkey, mouse, rat or pig.
- 15. (Previously presented) A method according to claim 11, wherein the pluripotent cells are embryonic stem cells, adult stem cells, mesenchymal stem cells, or umbilical cord blood cells.
  - 16-21. (Cancelled).